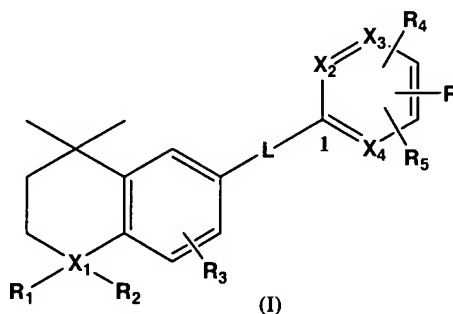


AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Claims 1-21. (Cancelled)

22. (new) A compound having a general formula (I):



wherein:

R is -C(O)NR₇R₈, -(CXY)_tC(O)NR₇R₈, -C(O)C(O)NHMe, -(C=C)C(O) NR₈R₉, -C(O)CF₃, or another Zn-chelating- group, with the proviso that R is not an acidic group or an ester derivative, -COOR₉ or salt thereof, R₇ is a group of formula -OH, -OR₉, 2-aminophenyl and R₈ is selected from hydrogen, C₁₋₆alkyl; R₉ is independently selected from hydrogen or C₁₋₆alkyl ; t is 1, 2 or 3 (preferably 1), X and Y, which are identical or different, represent an hydrogen or halogen atom (preferably F),

X₁ represents a carbon, oxygen, nitrogen or sulphur atom,

R₁ and R₂ represent independently or form together:

- . a C₁₋₆alkyl group, in particular methyl or ethyl groups, when X₁ is a carbon atom,
- . nothing, when X₁ is a oxygen or sulphur atom,
- . one or two oxygen atoms, when X₁ is sulphur atom (the case of a sulfoxide –SO- or a sulphone –SO₂-), or
- . one atom of hydrogen, an alkyl, aryl or aralkyl group, when X₁ is an atom of nitrogen (the case of an amino –NH, -an N-alkyl, N-aryl or N-aralkyl group);

X₂ and X₃, which are identical or different, represent CH, an atom of oxygen or an atom of nitrogen, or X₂=X₃ may be a single atom of sulphur, oxygen or nitrogen, or in the case where X₂ is an atom of oxygen and X₃ an atom of nitrogen, C₁ and X₄ represent a single one and same carbon atom, so that the ring carrying X₂ and X₃ can be an isoxazole ring,

X₄ can be CH or a nitrogen atom,

R₄ and R₅, which are identical or different, represent a hydrogen atom, a halogen atom, more particularly a fluorine atom, a C₁₋₆alkyl group, a group of formula -OH, –NH₂, -NHR₆, -OR₆, -SR₆, -(CF₂)_nCF₃, where n is an integer from 0 to 10, and whenever possible their salts with physiologically tolerated acids,

R₆ represents a hydrogen atom, a C₁₋₆alkyl group, a fluoroalkyl group having from 1 to 6 carbons atoms and from 3 to 7 fluorine atoms, an aryl group or an aralkyl group;

R₃ has the same definition as R₄ and R₅;

L is a linker and represents a bivalent radical either linear or cyclic, either saturated or unsaturated, more particularly L represents a bivalent radical derived from an alkane, alkene, alkyne or, aromatic or not, cyclic containing hydrocarbon group having from 1 to 12 carbon atoms, another bivalent radical of the following formula -O-, -CO-, -CO-NH-, -NH-CO-, -NH-CO-NH-, -CF₂-CO-NH-, -C(XY)-CO-NH-CH₂-, -NH-CO-CO-NH-, NH-CO-CO-NH-CH₂-, -SO₂NH-, -NH-SO₂-, -SO₂NCH₃-, -NCH₃SO₂-, -NR₆-, -C(=NOH)-, or a mixture thereof; R₆ being as defined above, optionally the bivalent radical is substituted, in particular by at least one C₁₋₆alkyl group;

X and Y, which are identical or different, represent an hydrogen or halogen atom (preferably F);

its tautomers, optical and geometrical isomers, racemates, salts, hydrates and mixtures thereof.

23. (new) A compound according to claim 22, wherein the groups identified in claim 1 are substituted with at least one substituent, which may be selected in the group consisting of: a hydrogen atom, a halogen atom (preferably F, Cl, or Br), a hydroxyl group, a C₁₋₁₀alkyl group, an alkenyl group, an C₁₋₁₀alkanoyl group, a (C₁-C₁₀)alkoxy group, a (C₁-C₁₀)alkoxycarbonyl group, an aryl group, an aralkyl group, an arylcarbonyl group, a mono- or poly-cyclic hydrocarbon group, a -NHCO(C₁-C₆)alkyl group, -NO₂, -CN, a -Nrr' group and a trifluoro(C₁-C₆)alkyl group, in which r and r', which are identical or different, represent a hydrogen atom, a lower alkyl group, an aryl, aralkyl group, an α -aminoacid group, a sugar group or a heterocycle group, or in which r and r' taken together form a heterocyclic ring.

24. (new) A compound according to claim 22, wherein R is -C(O)NR₇R₈ or - (CXY)_tC(O)NR₇R₈, in particular wherein R₈ is an hydrogen atom and R₇ is an hydroxyl group or a 2-aminophenyl group, preferably with X and Y are both halogen atoms and t is 1.

25. (new) A compound according to claim 22, wherein R is an hydroxamic acid group -(C=O)-NH-OH), a 2,2-difluoro-N-hydroxyacetamido group (-CF₂-(C=O)-NH-OH), or a N-(2-aminophenyl)acetamido group.

26. (new) A compound according to claim 22, wherein R is an electrophilic ketone, in particular -(C=O)-CF₃ or α -ketoamides, for instance -(C=O)-(C=O)-NHMe.

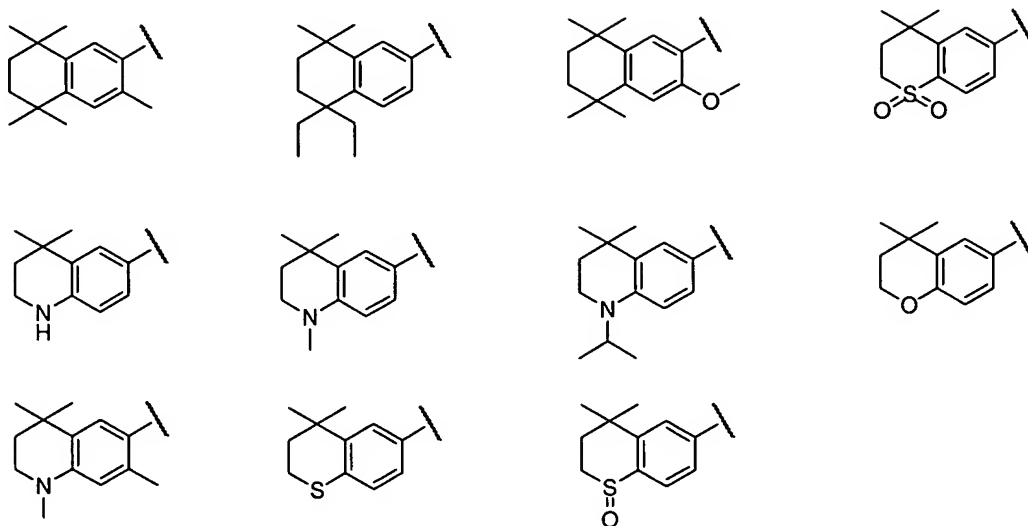
27. (new) A compound according to claim 22, wherein R is in para or meta position of C1, R is preferably in para position of C1.

28. (new) A compound according to claim 22, wherein L represents -CO-NH-, -NH-CO-, -CH=CH- (cis or trans forms), -CF₂-CO-NH-, -CF₂-CO-NH-CH₂-, or -NH-CO-CO-NH-.

29. (new) A compound according to claim 22, wherein R₃ is an hydrogen atom, OR₆, in particular methoxy, or a C₁₋₆alkyl group, in particular methyl.

30. (new) A compound according to claim 22, wherein R₃ is on position 2 of the substituted naphthalene derivative.

31. (new) A compound according to claim 22, wherein the ring carrying X₁ is selected from:



32. (new) A compound according to claim 22, wherein the ring carrying X_2 , X_3 , and X_4 is selected from phenyl, pyridinyl, pyrimidinyl, isoxazolyl, thiophenyl, furanyl, pyrrolyl, pyrazolyl, imidazolyl, isothiazolyl, thiazolyl, thienyl, thienooxazolyl and triazinyl rings.

33. (new) A compound according to claim 22, wherein the ring carrying X_2 , X_3 , and X_4 is phenyl, optionally substituted by a halogen atom, more particularly a fluorine atom, a C_{1-6} alkyl group, a group of formula $-OH$, or OR_6 .

34. (new) A compound, which is selected from the group consisting of:

N-(4-(Hydroxycarbamoyl)phenyl)-5,6,7,8-tetrahydro-5,5,8,8-tetramethylnaphthalene-2-carboxamide

N-(4-(2-Aminophenylcarbamoyl)phenyl)-5,6,7,8-tetrahydro-5,5,8,8-tetramethylnaphthalene-2-carboxamide

N-(1,2,3,4-Tetrahydro-1,1,4,4-tetramethylnaphthalen-6-yl)-*N'*-hydroxyterephthalamide

4-((*E*)-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-6-yl)vinyl)-*N*-hydroxybenzamide

4-((*Z*)-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-6-yl)vinyl)-*N*-hydroxybenzamide

4-(2,2-difluoro-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-7-yl)acetamido)-*N*-hydroxybenzamide

3-(2,2-difluoro-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-7-yl)acetamido)-*N*-hydroxybenzamide

4-((2,2-difluoro-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-7-yl)acetamido)methyl)-*N*-hydroxybenzamide

N-(4-((hydroxycarbamoyl)difluoromethyl)phenyl)-5,6,7,8-tetrahydro-5,5,8,8-tetramethylnaphthalene-2-carboxamide

N-(4-Hydroxycarbamoyl-phenyl)-*N'*-(5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-naphthalen-2-yl)-oxalamide

N-(4-Hydroxycarbamoyl-benzyl)-*N'*-(5,5,8,8-tetramethyl-5,6,7,8-tetrahydro-naphthalen-2-yl)-oxalamide.

35. (new) A compound, which is selected from the group consisting of:

4-(2,2-Difluoro-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-7-yl)acetamido)-*N*-hydroxybenzamide

N-(1,2,3,4-Tetrahydro-1,1,4,4-tetramethylnaphthalen-6-yl)-*N'*-hydroxyterephthalamide

4-((2,2-Difluoro-2-(1,2,3,4-tetrahydro-1,1,4,4-tetramethylnaphthalen-7-yl)acetamido)methyl)-*N*-hydroxybenzamide

N-(4-(Hydroxycarbamoyl)phenyl)-5,6,7,8-tetrahydro-5,5,8,8-tetramethylnaphthalene-2-carboxamide

36. (new) A pharmaceutical composition comprising at least one compound of formula (I), as defined in claim 22, and a pharmaceutically acceptable vehicle or support.

37. (new) A method for the treatment of conditions mediated by HDAC, such as cancers, in particular promyelocytic leukaemia, or other diseases associated with abnormal cell proliferation, such as psoriasis, comprising the administration to a subject in need thereof of an effective amount of a compound as defined in claim 22.

38. (new) A method for the treatment of central and peripheral nervous system diseases or neurodegenerative diseases associated with an excitotoxicity, such as Huntington's disease, such as polyglutamine expansion diseases, Alzheimer disease, Parkinson disease, multiple sclerosis, neuronal ischemia or amyotrophic lateral sclerosis (ALS), comprising the administration to a subject in need thereof of an effective amount of a compound as defined in claim 22.

39. (new) A method for the treatment of fibrosis, comprising the administration to a subject in need thereof of an effective amount of a compound as defined in claim 22.

40. (new) A method according to claim 37, wherein the cancer is selected from promyelocytic leukaemia, prostate cancer, ovarian cancer, pancreas cancer, lung cancer, breast cancer, liver cancer, head and neck cancer, colon cancer, bladder cancer, non-Hodgkin 's lymphoma cancer and melanoma.

41. (new) A method for reducing cancer cell proliferation, comprising the administration to a subject in need thereof of an effective amount of a compound as defined in claim 22.